25

Dye Mixtures Of Fibre-Reactive Azo Dyes And Use Thereof For Dyeing

Material Containing Hydroxy- And/Or Carboxamido Groups

The present invention relates to the field of fibre-reactive dyes.

Dyestuffs containing chromophores linked via a piperazine type linking unit are known from literature and are described for example in EP-A-0126265, EP-A-0693538, W099/05224 and W000/08104.

The inventors of the present invention have surprisingly found that mixtures of dyestuffs of the general formula (I) and dyestuffs of the general formula (II) give excellent application properties on cellulose containing material, especially high levels of solubility in water or salt solution, high fixation degrees, ease of washing out the unfixed dyestuff, good fastness to light and water as well as robustness to process variables.

The present invention claims mixtures of fibre reactive dyes comprising one or more dyestuffs of the formula (I)

and one or more dyestuffs of the general formula (II)

where

X¹, X² independently are a labile atom or group;

Ar¹ is an aromatic residue substituted by at least one -SO₃M group,
 M is hydrogen or alkali metal, especially sodium,

Ar² is an aromatic radical substituted with at least one -SO₃M group,

a is 1 or 2 wherein,

if a is 2

20

L is a divalent radical typically of the form (y)

where

15 R¹ and R² are independently hydrogen, C₁-C₄ alkyl optionally substituted by -OR, -SR, -SO₃M, or -X,

or a phenyl group optionally substituted by a sulfonic acid group, -OR, -C1- C4 alkyl, or - NR'COR

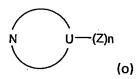
 L^1 is alkylene or arylene optionally substituted by a sulfonic acid group, -OR, -C $_1$ -C $_4$ alkyl or - NR'COR, -COOR wherein R and R' are independently hydrogen or C $_1$ - C $_4$ alkyl and X is halogen -

or R¹ and R² are independently optionally substituted alkyl, or is aminoethylpiperazine, under the proviso that if L is

25 aminopiperazine, Ar¹ and Ar² are different or

if a is 1

L is a monovalent radical -NR³R⁴, -SR³ or -OR³ where
R³ and R⁴ have one of the meanings of R¹ and R² or for -NR³R⁴, R³ and R⁴ can form a cyclic structure of the form (o)



where

10

U is a C_4 - C_6 alkyl residue optionally substituted by a substituent of formula Z and optionally interrupted by heteroatoms or heteroatom-containing groups such as - O-, - NR¹,

n is 1, 2 or 3 and

15

Z is hydrogen, optionally substituted C_1 - C_4 alkyl, -OR 5 , - CO_2R^5 , -COR 5 and

 R^5 is hydrogen, optionally substituted C_1 - C_4 alkyl, optionally substituted vinyl, optionally substituted phenyl.

20 (C₁-C₄)-alkyl groups may be straight-chain or branched and are preferably for example methyl, ethyl, n-propyl, i-propyl or n-butyl. Substituted alkyl groups are preferably substituted by hydroxyl, (C₁-C₄)-alkoxy, halogen or carboxyl groups. Substituted vinyl groups are for example – C(CH₃) = CH₂ or – CH = CHCOOH, substituted phenyl groups are for example phenyl substituted by – COOH, or – SO₃M.

Preferred embodiments of the invention are mixtures of one or more dyestuffs of the general formula (I) and one or more dyestuffs of the general formula (II)

30 where

 X^1 and X^2 are independently chlorine, fluorine or 3 or 4-carboxypyridinium especially preferred is chlorine,

Ar¹ and Ar² are independently a naphthyl residue substituted by at least one sulfo group (q-1)

where n is 1 to 3, especially preferred (q-11) or (q-12)

$$MO_3S$$
 or SO_3M $(q-11)$ $(q-12)$

10

15

5

or are a phenyl residue substituted by at least one sulfo group (q-2)

wherein

m is 1 or 2

p is 1 or 2

Y is independently hydrogen, halogen, R^5 , OR^5 , SR^5 , NHCOR 5 , where R^5 is as defined above, especially preferred Y is methyl.

When a is 2 t⁻¹ is preferred to be an optionally substituted phenylene or an alkylene residue optionally substituted or optionally interrupted by heteroatoms or heteroatom containing groups, wherein L¹-N-R¹ or L¹-N-R² may contain a cyclic structural feature such as

10

15

20

where n and R^1 are as defined above and x is 2 to 5 and Z' has one of the meanings of Z.

When a is 1 L is preferably morpholine or N-Methylsulfanilic acid.

The dyestuffs of the formula (I) are contained in the mixture in quantity of 1% by weight to 99% by weight preferably in a mixing ration of 10% by weight to 90% by weight and the dyestuffs of the formula (II) are contained in the mixture in a mixing ratio of 99% by weight to 1% by weight,

preferably in a mixing ratio of 90% by weight to 10% by weight.

Dyestuff mixtures according to the invention can be obtained upon reacting chromophores of formula (III)

with an appropriate mixture of 2-aminoethylpiperazine and a diamine H-L-H, or amine H-L, wherein L is as defined above, followed by precipitation using methylated spirits and conventional filtration.

The dyestuffs of the present invention can be present as a preparation in solid or liquid (dissolved) form. In solid form they generally contain the electrolyte salts customary in the case of water-soluble and in particular fibre-reactive dyes, such as sodium chloride, potassium chloride and sodium

sulfate, and also the auxiliaries customary in commercial dyes, such as buffer substances capable of establishing a pH in aqueous solution between 3 and 8, such as sodium acetate, sodium borate, sodium bicarbonate, sodium citrate, sodium dihydrogenphosphate and disodium

- hydrogenphosphate, small amounts of siccatives or, if they are present in liquid, aqueous solution (including the presence of thickeners of the type customary in print pastes), substances which ensure the permanence of these preparations, for example mold preventatives.
- In general, the dyestuff mixtures of the present invention are present as dye 10 powders containing 10 to 80% by weight, based on the dye powder or preparation, of a strength-standardizing colorless diluent electrolyte salt, such as those mentioned above. These dye powders may additionally include the aforementioned buffer substances in a total amount of up to 10%, based on the dye powder. If the dye mixtures of the present invention are present 15 in aqueous solution, the total dye content of these aqueous solutions is up to about 50 % by weight, for example between 5 and 50% by weight, and the electrolyte salt content of these aqueous solutions will preferably be below 10% by weight, based on the aqueous solutions. The aqueous solutions (liquid preparations) may include the aforementioned buffer substances in an 20 amount which is generally up to 10% by weight, for example 0.1 to 10% by weight, preference being given to up to 4% by weight, especially 2 to 4% by weight.
- The dyestuff mixtures of the instant invention are reactive dyestuffs suitable for dyeing and printing hydroxy- and/or carboxamido-containing fibre materials by the application and fixing methods numerously described in the art for fibre-reactive dyes. They provide exceptionally strong and economic shades. Such dyes especially when used for exhaust dyeing of cellulosic materials can exhibit excellent properties including build-up, light-fastness, high levels of solubility in water or salt solution, high fixation degrees, ease of washing out the unfixed dyestuff, as well as robustness to process

variables.

10

15

30

They are also wholly compatible with similar dyes designed for high temperature (80-100°C) application to cellulosic textiles, and thus lead to highly reproducible application processes, with short application times. The present invention therefore also provides for use of the inventive dyestuffs for dyeing and printing hydroxy- and/or carboxamido-containing fibre materials and processes for dyeing and printing such materials using a dyestuff according to the invention. Usually the dyestuff is applied to the substrate in dissolved form and fixed on the fibre by the action of an alkali or by heating or both.

Hydroxy-containing materials are natural or synthetic hydroxy-containing materials, for example cellulose fiber materials, including in the form of paper, or their regenerated products and polyvinyl alcohols. Cellulose fiber materials are preferably cotton but also other natural vegetable fibers, such as linen, hemp, jute and ramie fibres. Regenerated cellulose fibers are for example staple viscose and filament viscose.

- Carboxamido-containing materials are for example synthetic and natural polyamides and polyurethanes, in particular in the form of fibers, for example wool and other animal hairs, silk, leather, nylon-6,6, nylon-6, nylon-11, and nylon-4.
- Application of the inventive dyestuffs is by generally known processes for dyeing and printing fiber materials by the known application techniques for fibre-reactive dyes. The dyestuffs according to the invention are highly compatible with similar dyes designed for high temperature (80-100°C) applications and are advantageously useful in exhaust dyeing processes.

Similarly, the conventional printing processes for cellulose fibers, which can either be carried out in single-phase, for example by printing with a print

15

20

25

30

paste containing sodium bicarbonate or some other acid-binding agent and the colorant, and subsequent steaming at appropriate temperatures, or in two phases, for example by printing with a neutral or weakly acid print paste containing the colorant and subsequent fixation either by passing the printed material through a hot electrolyte-containing alkaline bath or by overpadding with an alkaline electrolyte-containing padding liquor and subsequent batching of this treated material or subsequent steaming or subsequent treatment with dry heat, produce strong prints with well defined contours and a clear white ground. Changing fixing conditions has only little effect on the outcome of the prints. Not only in dyeing but also in printing the degrees of fixation obtained with dye mixtures of the invention are very high. The hot air used in dry heat fixing by the customary thermofix processes has a temperature of from 120 to 200°C. In addition to the customary steam at from 101 to 103°C, it is also possible to use superheated steam and high pressure steam at up to 160°C.

The inventive dyestuffs can in addition be used to produce inks useful for printing the substrates described above, for example textiles, especially cellulosic textiles, and paper. Such inks can be used in all technologies, for example conventional printing, ink-jet printing or bubble-jet printing (for information on such printing technologies see for example Text. Chem. Color, Volume 19(8), pages 23 ff and Volume 21, pages 27 ff).

Acid-binding agents responsible for fixing the dyes to cellulose fibers are for example water-soluble basic salts of alkali metals and of alkaline earth metals of inorganic or organic acids, and compounds, which release alkali when hot. Of particular suitability are the alkali metal hydroxides and alkali metal salts of weak to medium inorganic or organic acids; the preferred alkali metal compounds being the sodium and potassium compounds. These acid-binding agents are for example sodium hydroxide, potassium hydroxide, sodium carbonate, sodium bicarbonate, potassium carbonate, sodium formate, sodium dihydrogenphosphate and disodium hydrogenphosphate.

30

Treating the dyestuffs according to the invention with the acid-binding agents, with or without heating, bonds the dyes chemically to the cellulose fibers. Especially the dyeings on cellulose, after they have been given the usual aftertreatment of rinsing to remove unfixed dye portions, show excellent properties.

The dyeings of polyurethane and polyamide fibers are customarily carried out from an acid medium. The dyebath may contain for example acetic acid and/or ammonium sulfate and/or acetic acid and ammonium acetate or 10 sodium acetate to bring it to the desired pH. To obtain a dyeing of acceptable levelness it is advisable to add customary leveling auxiliaries, for example based on a reaction product of cyanuric chloride with three times the molar amount of an aminobenzenesulfonic acid or 15 aminonaphthalenesulfonic acid or based on a reaction product of for example stearylamine with ethylene oxide. In general the material to be dyed is introduced into the bath at a temperature of about 40°C and agitated therein for some time, the dyebath is then adjusted to the desired weakly acid, preferably weakly acetic acid, pH, and the actual dyeing is carried out at temperature between 60 and 98°C. However, the dyeings can also be 20 carried out at the boil or at temperatures up to 120°C (under superatmospheric pressure).

The examples hereinbelow serve to illustrate the invention. Parts and percentages are by weight, unless otherwise stated. Parts by weight relate to parts by volume as the kilogram relates to the liter. The compounds described in the examples in terms of a formula are indicated in the form of the free sulphonic acids, but as in general they are prepared and isolated in the form of their alkali metal salts, such as lithium, sodium or potassium salts, and used for dyeing in the form of these salts. The starting compounds and components mentioned in the form of the free acid in the examples hereinbelow may be used in the synthesis as such or similarly in the form of

their salts, preferably alkali metal salts.

5 Example 1

A 1:1 molar mixture of 2-aminoethylpiperazine (4) and ethylenediamine (5) (0.006 mol) was added to a stirred suspension of the dichlorotriazinyl red dye(3) (0.012 mol) in water (350 mols) at pH 6. The pH was adjusted and maintained at pH 10 by the addition of 2N NaOH solution. After three hours, chromatography indicated that the reaction was complete and the pH was adjusted to 6 with 2N HCl before precipitating the product by addition of methylated spirits. The resulting solid was filtered off and dried to give a dark red powder (11.3g). Analytical data were consistent with a 1:1 mixture of the two dyes (6) and (7).

15

10

The examples herein below describe further dye mixtures according to the invention. They may be prepared according to the invention either by mechanically mixing the individual dyes or else chemically, for example similarly to the above illustrative embodiment, using as starting compounds of general formula (III), in which X is for example chlorine and Ar¹ and Ar² are for example:

$$SO_3M$$
 SO_3M
 SO_3

10

2-aminopiperazine and a diamine H-L-H or amine H-L, wherein L is for example a residue of formula a to af

Following exactly analogous procedures the following dyes (examples 2 – 56) were synthesized giving 1:1 mixtures of the respective dyestuff (8) and (9)

Example No.	Ar ¹	Ar ²	. Ľ
2	Α	Α	а
3	Α	Α	°b
4	A	Α	С
5 .	Α .	A	d
6	Α	A	е
7	Α	A;	f
8	Α	Α	g
9	Α	Α	, h
10	Α	Α .	k
11	Α	Α	1

(9)

Example No.	Ar¹	Ar ²	L
12	Α	A	p
13	В	В	. а
14	В	В	d
15	В	В	g
16	В	В	k
17	В	В	1 .
18	В	В	.· m
19	В	В	o
20	В	В	р
21	Α	G	j
22	Α	G	k
23	Α	G	1.
24	В	F	k
25	В	F	!
26	В	F	0
27	В	F	р
28	С	С	а
29	С	С	k
30	С	С	- 1
31	F	F .	а
32	F	F	k
33	F	F	1
34	. D	D	k
35	D .	D	0
36	E	Ε	1
37	Н	Н	m
38	Н	Н	n
39	G	G	1

Ar ¹	Ar²	L
G	G	n,
В	В	V
F	F	· v
В	В	· w
F	F .	w
В	В	x
F	F	x
В	В	f
F	F	У
F	F	Z
В	В	aa
В	В	j
F	F	j
F	F	O
Α	Α	ab
В	В	ae
В .	В	af
	G B F B F B F F A B	G G B B F F B B F F B B F F F F B B B F F F F

Examples 57-71 consist of mixtures of dyes of the form (8) and (10).

	1	<u> </u>	
Example No.	Ar ¹	Ar ²	L
57	Α	A	q
58	Α	Α	r
59	Α	А	u
60	Α	Α .	t
61	В	В	. q
62	В	В	S
63	В	В	t
64	В.	В	u
65	F	F	q
66	F	F	S
67	F	F	t
68	С	С	S
69	С	С	t
70	G	G	u
71	В	В	ad

All these dyestuff mixtures give excellent application properties on cellulose containing material, especially high levels of solubility in water or salt solution, high fixation degrees, ease of washing out the unfixed dyestuff, good fastness to light and water as well as robustness to process variables.

What is claimed is:

1. Mixture of fibre reactive dyes comprising one or more dyestuffs of the formula (I)

5

and one or more dyestuffs of the general formula (II)

10

where

 X^1 , X^2 are independently a labile atom or group;

Ar¹ is an aromatic residue substituted by at least one -SO₃M group;

M is hydrogen or alkali metal, especially sodium;

15 Ar² is an aromatic radical substituted with at least one -SO₃M group;

a is 1 or 2 wherein, if a is 2

L is a divalent radical typically of the form (y)

where

 $_{\rm t}$ R¹ and R² are independently hydrogen, C₁-C₄ alkyl optionally substituted by – OR, -SR, -SO₃M or X, or a phenyl group optionally substituted by a sulfonic acid group, -OR, -C₁-C₄- alkyl, or NR'COR and L¹ is arylene or alkylene optionally substituted by a sulfonic acid group, -OR, -C₁-C₄-alkyl – COOR, or NR'COR or

10

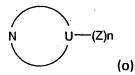
15

L is aminoethylpiperazine, under the proviso that if L is aminopiperazine, Ar¹ and Ar² are different or

if a is 1

L is a monovalent radical -NR³R⁴, -SR³ or -OR³ where

R³ and R⁴ have one of the meanings of R¹ and R² or for -- NR³R⁴, R³ and R⁴ can form a cyclic structure of the form (o)



20

25

where

U is an C_4 - C_6 alkyl residue optionally substituted by a substituent of formula Z and optionally interrupted by heteroatoms or heteroatom-containing groups such as - O-, $-NR^1$, n is 1, 2 or 3 and Z is hydrogen, optionally substituted C_1 - C_4 alkyl, $-OR^5$, $-CO_2R^5$, $-COR^5$

and

 R^5 is hydrogen, optionally substituted C_1 - C_4 alkyl, optionally substituted vinyl, optionally substituted phenyl.

2. Dyestuff mixture according to claim 1

5 wherein

X¹ and X² is independently chlorine, fluorine or 3 or 4-carboxypyridinium; Ar¹ and Ar² is independently a naphthyl residue substituted by at least one sulfo group (q-1)

where n is 1 to 3

or is a phenyl residue substituted by at least_one sulfo group (q-2)

(MO₃S)m

*
(Y)p (q-2)

15 wherein

m is 1 or 2

p is 1 or 2 and

Y is independently hydrogen, halogen, R^5 , OR^5 , SR^5 , $NHCOR^5$, where R^5 is as given in claim 1.

20

25

10

3. Dyestuff mixture according to claim 1

wherein

X¹ and X² is chlorine;

Ar¹ and Ar² are independently a naphthyl residue of the formula (q-11) or (q-12)

$$MO_3$$
S

or

 SO_3M
 SO_3M
 $(q-11)$
 $(q-12)$

or are a phenyl residue substituted by at least one sulfo group (q-2)

wherein

m is 1 or 2

p is 1 or 2 and

Y is methyl.

10

15

20

5

- 4. Dyestuff mixture according to claim 1 wherein a is 1 and L is morpholine.
- 5.A dye mixture according to at least one of the claims 1 to 4 wherein a dye of formula (I) is present in the mixture in an amount of from 1% by weight to 99% by weight and a dye of the formula (II) is present in the mixture in an amount of from 99% by weight to 1% by weight.
- 6. A dye mixture according to at least one of the claims 1 to 4 wherein a dye of formula (I) is present in the mixture in an amount of from 10% by weight to 90% by weight and a dye of the formula (II) is present in the mixture in an amount of from 90% by weight to 10% by weight.
- 7. A process for preparing a dye mixture as claimed in one or more of claims 1 to 5, which comprises reacting chromophores of formula (III)

. 5

10

wherein Ar¹, Ar², X¹, X² and M are as defined in claim 1 with an appropriate mixture of 2-aminoethylpiperazine and a diamine H-L-H, or amine H-L, wherein L is as defined above, followed by precipitation using methylated spirits and conventional filtration.

8. A process for dyeing hydroxy- and/or carboxamido – containing fiber material, in which dyestuffs or dyestuff mixtures are applied to the material and the dyes are fixed to the material by means of heat or with the aid of an alkali or by means of heat and with the aid of an alkali, which comprises dye mixtures or dyestuffs as claimed in one or more of the claims 1 to 5.

	1	PCT/E	3/12271
A. CLASS IPC 7	SIFICATION OF SUBJECT MATIL C09B67/00		
According	to International Patent Classification (IPC) or to both national classification and IPC		
	SSEARCHED		
Minimum of IPC 7	documentation searched (classification system followed by classification symbols) ${\tt C09B}$		·
Document	ation searched other than minimum documentation to the extent that such documents are includ	ed in the field	s searched
Electronic	data base consulted during the international search (name of data base and, where practical, s	earch terms us	sed)
	nternal, CHEM ABS Data		
C. DOCUM	ENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the retevant passages		Relevant to claim No.
Y	EP 0 693 538 A (CIBA GEIGY AG) 24 January 1996 (1996-01-24) cited in the application claim 1		1-8
Y	US 5 892 006 A (TZIKAS ATHANASSIOS) 6 April 1999 (1999-04-06) claim 1		1-8
Y	DE 100 64 496 A (DYSTAR TEXTILFARBEN GMBH & CO) 4 July 2002 (2002-07-04) claim 1		1-8
A	WO 99 05244 A (CONNOR DANIEL STEDMAN; SCHEIBEL JEFFREY JOHN (US); VINSON PHILLIP) 4 February 1999 (1999-02-04) cited in the application claim 1		1-8
<u> </u>	er documents are listed in the continuation of box C.	mbers are liste	d in annex.
Special cat	egories of cited documents :	od after the '-	Introdica al Cita and the
conside	nt defining the general state of the art which is not clied to understand the pred to be of particular relevance clied to understand the ocument but published on or after the international	it in conflict wit e principle or t	h the application but

Further documents are listed in the continuation of box C.	X Palent family members are listed in annex.
Special categories of cited documents: A document defining the general state of the art which is not considered to be of particular relevance E earlier document but published on or after the international filing date L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) O document referring to an oral disclosure, use, exhibition or other means P document published prior to the international filing date but later than the priority date claimed	 "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. "&" document member of the same patent family
Date of the actual completion of the international search	Date of mailing of the international search report
25 February 2004	08/03/2004
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk	Authorized officer
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Bakboord, J

Form PCT/ISA/210 (second sheet) (July 1992)

ppiication No

T/LP U3/12271

Category °	ation) DOCUMENTS CONSIDERED TO BE RELEVANT Citation of document, with indication, where appropriate, of the relevant passages	
Juligoty	ordinent, with intucation, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 00 08104 A (BASF AG ;EBENEZER WARREN JAMES (GB); MYNETT DONNA MARIA (GB)) 17 February 2000 (2000-02-17) cited in the application claim 1	1-8
P,A	WO 02 092697 A (DYSTAR TEXTILFARBEN GMBH & CO ;EBENEZER WARREN JAMES (GB)) 21 November 2002 (2002-11-21) claim 1	1-8
P,A	EP 1 380 621 A (DYSTAR TEXTILFARBEN GMBH & CO) 14 January 2004 (2004-01-14) claim 1	1-8
	. •	
	•	
,		
	•	

	ation on patent family me	embers	PCT/t	3/12271
Patent document cited in search report	Publication date		Patent family member(s)	Publication date
EP 0693538 A	24-01-1996	BR CN DE EP ES GR JP PT US	9502861 A 1133323 A ,B 59509529 D1 0693538 A2 2161847 T3 3036651 T3 8003469 A 693538 T 5631352 A	04-06-1996 16-10-1996 27-09-2001 24-01-1996 16-12-2001 31-12-2001 09-01-1996 30-01-2002 20-05-1997
US 5892006 A	06-04-1999	US US DE DE EP EP ES ES HK JP JP JP US	5735911 A 5612463 A 5451665 A 59108779 D1 59109196 D1 59109201 D1 0478503 A2 0735107 A2 0735113 A2 2106070 T3 2150620 T3 2152455 T3 1001093 A1 3231364 B2 4258674 A 3389555 B2 2001026726 A 2002194241 A 5232462 A	07-04-1998 18-03-1997 19-09-1995 21-08-1997 05-10-2000 23-11-2000 01-04-1992 02-10-1996 01-11-1997 01-12-2000 01-02-2001 22-05-1998 19-11-2001 14-09-1992 24-03-2003 30-01-2001 10-07-2002 03-08-1993
DE 10064496 A	04-07-2002	DE BR CA WO EP	10064496 A1 0116419 A 2428054 A1 02051944 A1 1345994 A1	04-07-2002 30-12-2003 05-05-2003 04-07-2002 24-09-2003
WO 9905244 A	04-02-1999	AU AU BR CA CN EP HU WO ID JP TR US US ZA	738353 B2 8124998 A 9811524 A 2297171 C 1270622 T 1002031 A1 0002626 A2 9905244 A1 28301 A 2001511474 T 200000923 T2 2002103096 A1 6306817 B1 9806447 A	13-09-2001 16-02-1999 18-12-2001 01-04-2003 18-10-2000 24-05-2000 28-11-2000 04-02-1999 10-05-2001 14-08-2001 21-09-2000 01-08-2002 23-10-2001 21-01-1999
WO 0008104 A	17-02-2000	AT BR DE DE EP ES WO	237661 T 9912628 A 69906983 D1 69906983 T2 1100847 A1 2197658 T3 0008104 A1	15-05-2003 02-05-2001 22-05-2003 18-12-2003 23-05-2001 01-01-2004 17-02-2000

tion on	patent fan	nily members
---------	------------	--------------

i	<u>:</u>			Į.	J/EP	03/12271
Patent document cited in search report		Publication date		Patent family member(s)		Publication date
WO 0008104	A		JP PT TR US	2002522587 1100847 200100320 6359121	T T2	23-07-2002 31-07-2003 21-06-2001 19-03-2002
WO 02092697	Α	21-11-2002	CA WO EP	2446784 02092697 1387865	A1	21-11-2002 21-11-2002 11-02-2004
EP 1380621	A	14-01-2004	CA EP	2434825 1380621		10-01-2004 14-01-2004

, interna

pplication No

This Page is Inserted by IFW Indexing and Scanning Operations and is not part of the Official Record

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

☐ BLACK BORDERS	• .	
☑ IMAGE CUT OFF AT TOP, BOTTOM OR SIDES		
☐ FADED TEXT OR DRAWING		
☐ BLURRED OR ILLEGIBLE TEXT OR DRAWING		
SKEWED/SLANTED IMAGES		
☐ COLOR OR BLACK AND WHITE PHOTOGRAPHS		
☐ GRAY SCALE DOCUMENTS		
☐ LINES OR MARKS ON ORIGINAL DOCUMENT		
☐ REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUAL	ITY	
Потиер.		

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.